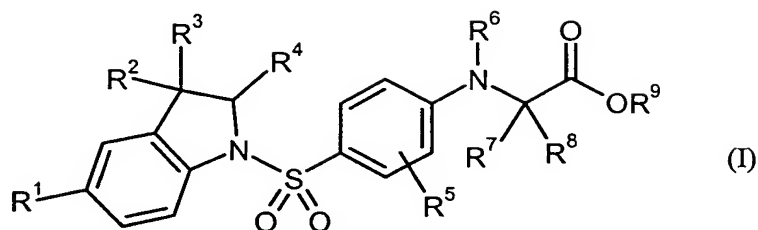


Patent Claims -

1. Compounds of the general formula (I)



in which

- 5 R^1 represents phenyl or represents 5- or 6-membered heteroaryl having up to two heteroatoms from the group consisting of N, O and S, which radicals may for their part each be mono- to trisubstituted by identical or different substituents selected from the group consisting of halogen, cyano, nitro, (C₁-C₆)-alkyl (which for its part may be substituted by hydroxyl), (C₁-C₆)-alkoxy, trifluoromethyl, trifluoromethoxy, (C₁-C₆)-alkylsulphonyl, (C₁-C₆)-alkanoyl, (C₁-C₆)-alkoxycarbonyl, carboxyl, amino, (C₁-C₆)-acylamino, mono- and di-(C₁-C₆)-alkylamino,

- 15 R^2 and R^3 are identical or different and independently of one another represent hydrogen or (C₁-C₄)-alkyl or together with the carbon atom to which they are attached form a 3- to 7-membered spiro-linked cycloalkyl ring,

R^4 represents hydrogen or (C₁-C₄)-alkyl,

R^5 represents hydrogen, (C₁-C₄)-alkyl, (C₁-C₄)-alkoxy or halogen,

R^6 represents (C₁-C₆)-alkyl, (C₃-C₈)-cycloalkyl, (C₁-C₆)-alkanoyl, (C₁-C₆)-alkylsulphonyl or (C₁-C₆)-alkoxycarbonyl,

- 20 R^7 and R^8 are identical or different and independently of one another represent hydrogen or (C₁-C₄)-alkyl,

and

R^9 represents hydrogen or a hydrolyzable group which can be degraded to the corresponding carboxylic acid,

and their pharmaceutically acceptable salts, solvates and solvates of the salts.

2. Compounds of the general formula (I) according to Claim 1 in which

R¹ represents phenyl which may be mono- or disubstituted by identical or different substituents selected from the group consisting of fluorine, chlorine, cyano, (C₁-C₄)-alkyl, (C₁-C₄)-alkoxy, trifluoromethyl, trifluoromethoxy, methylsulphonyl, acetyl, propionyl, (C₁-C₄)-alkoxycarbonyl, amino, acetylamino, mono- and di-(C₁-C₄)-alkylamino,

R² and R³ are identical or different and independently of one another represent hydrogen or (C₁-C₄)-alkyl or together with the carbon atom to which they are attached form a 5- or 6-membered spiro-linked cycloalkyl ring,

R⁴ represents hydrogen or methyl,

R⁵ represents hydrogen, methyl, methoxy, fluorine or chlorine,

R⁶ represents (C₁-C₄)-alkyl, acetyl, methylsulphonyl, methoxycarbonyl or tert-butoxycarbonyl,

R⁷ and R⁸ are identical or different and independently of one another represent hydrogen or methyl,

and

R⁹ represents hydrogen.

3. Compounds of the general formula (I) according to Claim 1 in which

R¹ represents phenyl which may be mono- or disubstituted by identical or different substituents selected from the group consisting of fluorine, chlorine, methyl, trifluoromethyl and trifluoromethoxy,

R² represents methyl,

R³ represents methyl,

or

R² and R³ together with the carbon atom to which they are attached form a spiro-linked cyclopentane or cyclohexane ring,

R^4 represents hydrogen or methyl,

R^5 represents hydrogen, methyl, fluorine or chlorine,

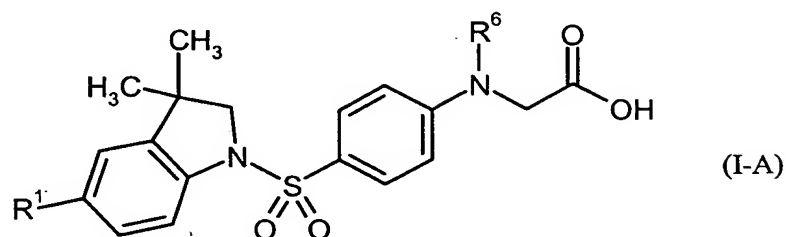
R^6 represents (C₁-C₄)-alkyl, acetyl or methylsulphonyl,

R^7 and R^8 each represent hydrogen

5 and

R^9 represents hydrogen.

4. Compounds of the formula (I-A)



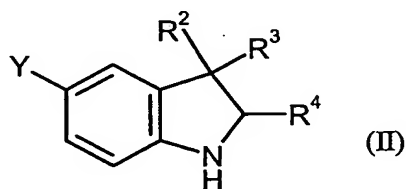
in which

10 R^1 represents phenyl which is substituted by fluorine, chlorine or trifluoromethyl,

and

R^6 represents methyl, ethyl, n-propyl, isopropyl or tert-butyl.

5. Process for preparing the compounds of the general formula (I) or (I-A) as defined in Claims 1 to 4, characterized in that compounds of the formula (II)



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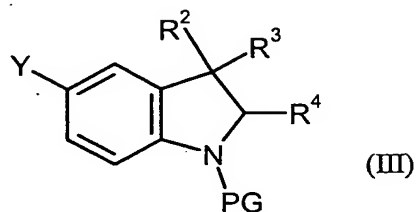
in which

R^2 , R^3 and R^4 are each as defined in Claim 1

and

Y represents chlorine or bromine

are initially, by methods known from the literature, converted into compounds of the formula (III)



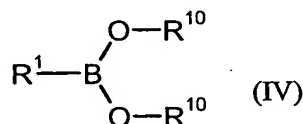
5 in which

Y, R², R³ and R⁴ are each as defined in Claim 1

and

PG represents a suitable amino protective group, preferably 4-nitrophenylsulphonyl,

10 these compounds are then reacted in a coupling reaction with a compound of the formula (IV)



in which

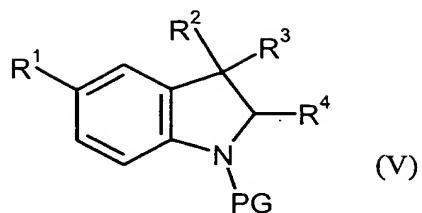
R¹ is as defined in Claim 1

and

15 R¹⁰ represents hydrogen or methyl or both radicals together form a CH₂CH₂- or C(CH₃)₂-C(CH₃)₂-bridge

in an inert solvent in the presence of a suitable palladium catalyst and a base to give compounds of the formula (V)

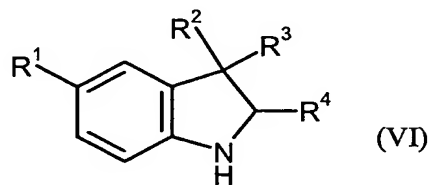
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in which

PG, R¹, R², R³ and R⁴ are each as defined in Claim 1,

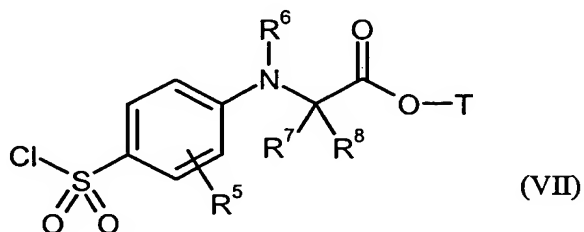
the protective group PG is then, by methods known from the literature, removed again
5 giving compounds of the formula (VI)



in which

R¹, R², R³ and R⁴ are each as defined in Claim 1,

the product is then converted with a compound of the formula (VII)



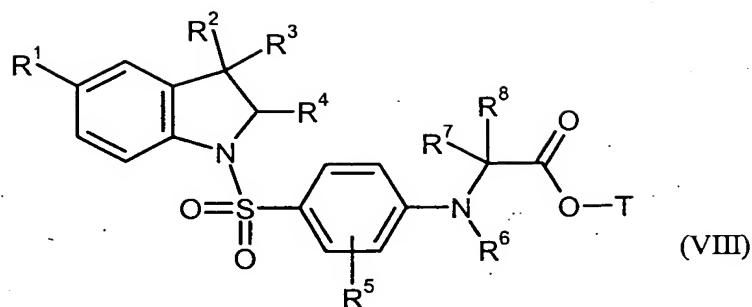
in which

R⁵, R⁶, R⁷ and R⁸ are each as defined in Claim 1

and

T represents benzyl or (C₁-C₆)-alkyl

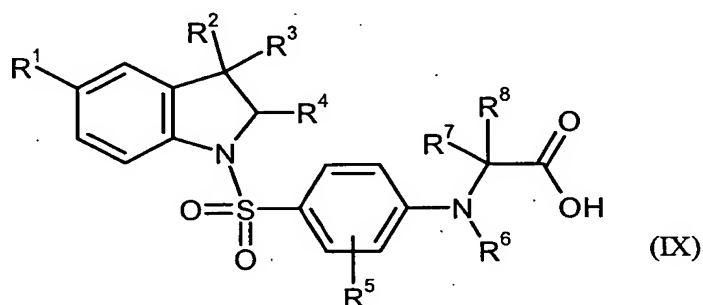
in an inert solvent in the presence of a base into compounds of the formula (VIII)



in which

T, R¹, R², R³, R⁴, R⁵, R⁶, R⁷ and R⁸ are each as defined in Claim 1,

the compounds of the formula (VIII) are then, using acids or bases or, if T represents benzyl, also hydrogenolytically, converted into the corresponding carboxylic acids of the formula (IX)



in which

R¹, R², R³, R⁴, R⁵, R⁶, R⁷ and R⁸ are each as defined in Claim 1,

these carboxylic acids (IX) are, if appropriate, modified further by known esterification methods to give compounds of the formula (I),

and the resulting compounds of the formula (IX) or (I) are, if appropriate, converted into their solvates, salts and/or solvates of the salts using the corresponding (i) solvents and/or (ii) bases or acids.

6. Compounds of the formula (I) or (I-A) as defined in Claims 1 to 4 for the prophylaxis and treatment of diseases.

7. Medicaments, comprising at least one compound of the formula (I) or (I-A) as defined in Claims 1 to 4 and inert non-toxic pharmaceutically acceptable carriers, auxiliaries, solvents, vehicles, emulsifiers and/or dispersants.
8. Use of compounds of the formula (I) or (I-A) and medicaments as defined in Claims 1 to 7
5 for the prophylaxis and treatment of diseases.
9. Use of compounds of the formula (I) or (I-A) as defined in Claims 1 to 6 for preparing medicaments.
10. Use of compounds of the formula (I) or (I-A) as defined in Claims 1 to 4 for preparing medicaments for the prophylaxis and treatment of stroke, arteriosclerosis, coronary heart
10 diseases and dyslipidaemias, for the prophylaxis of myocardial infarction and for the treatment of restenosis after coronary angioplasty or stenting.
11. Method for the prophylaxis and treatment of diseases, characterized in that compounds of the formula (I) or (I-A) as defined in Claims 1 to 4 are allowed to act on living beings.